Claims

[1] 1. A compound of the following formula (1):

(1)

in which

I) R represents H, simple alkyl chain (-SAC), simple cycloalkyl chain (-SCAC), aryl group (-Ar), or simple alkyl chain substituted by aryl (-SAC-Ar),

II) R¹ represents -SAC, -SCAC, -Ar, -SAC-Ar, or a side chain residue of all the natural amino acids; and the compound of formula (1) may exist in a specific diastereomeric form, or mixtures thereof when the carbon to which R¹ is attached becomes a stereocenter due to the R¹ group; or the compound of formula (1) may have a protecting group in an ester form (-CO R³ wherein R³ is -SAC) or a sulfonamide form (-CONHSO R⁴ wherein R⁴ is -SAC), or may exist in the form of pharmaceutically acceptable salt, when R¹ is a side chain residue of an amino acid containing carboxyl moiety; or the compound of formula (1) may also exist in the form of pharmaceutically acceptable salt when R¹ is a side chain residue of an amino acid containing a base moiety,

III) R² represents -SAC, -SCAC, -Ar, -SAC-Ar, or a side chain residue of the natural amino acids; and the compound of formula (1) may exist in a specific diastereomeric form, or mixtures thereof when the carbon to which R² is attached becomes a stereocenter due to the R² group; the compound of formula (1) may have a protecting group in an ester form (-CO R⁵ wherein R⁵ is -SAC) or a sulfonamide form (-CONHSO R⁶ wherein R⁶ is -SAC), or may exist in the form of pharmaceutically acceptable salt, when R² is a side chain residue of an amino acid containing carboxyl moiety; or the compound of formula (1) may also exist in the form of pharmaceutically acceptable salt when R² is a side chain residue of an amino acid containing a base moiety, or

 R^2 further represents H; -(CH₂) OR⁷ wherein R⁷ is -SAC, -SCAC, -Ar, or - SAC-Ar, and n = 1 or 2; or -(CH₂) OC(=O)R⁸ wherein R⁸ is -SAC, -SCAC, -Ar, or - SAC-Ar, and n = 1 or 2,

IV) A represents -(CH₂) - (n = 0-4), -O-(CH₂) - (n = 0-4), or -NR⁹-(CH₂) - (n = 0-4) wherein R⁹ is -SAC, -SCAC, -Ar, or -SAC-Ar,

V) Brepresents H, -SAC, -SCAC, -Ar, or -SAC-Ar, or

VI) R and R¹ may form a cycle together with the carbon atom to which they are attached, where -R-R¹- is -(CH₂) -, -(CH₂) -O-(CH₂) -, or -(CH₂) -NR¹⁰-(CH₂) - wherein n+m<9 and R¹⁰ is -SAC, -SCAC, -Ar, -SAC-Ar, -C(=O)-SAC, -C(=O)-SAC, -C(=O)-Ar, or -C(=O)-SAC-Ar,

VII) X represents -C(=O)CH OR¹¹ wherein R¹¹ is -SAC, -SCAC, -Ar, or -SAC-Ar; -C(=O)CH OC(=O)R¹² wherein R¹² is -SAC, -SCAC, -Ar, or -SAC-Ar; -CH=CH-CO R¹³ wherein R¹³ is -SAC, -SCAC, -Ar, or -SAC-Ar; -CH=CH-SO R¹⁴ wherein R¹⁴ is -SAC, -SCAC, -Ar, or -SAC-Ar; -C(=O)CH=CH₂; or -COCH₂-W wherein W is -N₂, -F, -Cl₂, -Br, -I, -NR¹⁵R¹⁶ (R¹⁵ and R¹⁶ each are -SAC, -SCAC, -Ar, or -SAC-Ar, or together may form 3- to 6-membered saturated or unsaturated cyclic group), -SR¹⁷ (R¹⁷ is -SAC, -SCAC, -Ar, or -SAC-Ar), or is the following formula:

wherein

Y is H, -OH, -OR¹⁸ (R¹⁸ = -SAC or -SCAC), -C(=O)R¹⁹ (R¹⁹ = -H, -SAC, or -SCAC), -F, -Cl, -Br, -I, -CN, -NC, -N₃, -CO₂H, CF₃, -CO₂R²⁰ (R²⁰ = -SAC or -SCAC), -C(=O)NHR²¹ (R²¹ = -SAC or -SCAC), or -C(=O)NR²²R²³ (R²² and R²³ each are -SAC, -SCAC, -Ar, or -SAC-Ar), R²⁴ is H, -SAC, -SAC-Ar, or -Ar, salt, or stereoisomer thereof.

O Tile annual according to plaim 1 wherein D represents U

- [4] 2. The compound according to claim 1 wherein R represents H.
- [5] 3. The compound according to claim 1 wherein R^1 represents -CH₂COOH, -CH₂COOH, or -CH₂CONHSO₂ R^4 (R^4 = SAC).
- 4. The compound according to claim 1 wherein R² represents H, -SAC, -Ar, or (CH₂) OR⁷ (R⁷ = -SAC, -SCAC, -Ar, or -SAC-Ar, and n = 1 or 2).
- 5. The compound according to claim 1 wherein X represents -C(=O)CH OAr, -C(=O)CH OC(=O)Ar, or -COCH -W wherein W is -N, -F, -Cl, -Br, -I, -NR R R and R -SAC, -SCAC, -Ar, or -SAC-Ar, or together may form 3- to 6-membered saturated or unsaturated cyclic group), or -SR R SCAC, -Ar, or -SAC-Ar).
- [8] 6. The compound according to claim 1 whereinI) R represents H,

```
II) R^1 represents -CH<sub>2</sub>COOH, -CH<sub>2</sub>COOR<sup>3</sup> (R^3=SAC), or -CH<sub>2</sub>CONHSO<sub>2</sub>R^4 (R^4 = SAC),
```

- III) R^2 represents H, -SAC, -Ar, or -(CH₂) OR^7 (R^7 = -SAC, -SCAC, -Ar, or -SAC-Ar, and n = 1 or 2),
- IV) A represents -(CH₂)_n (n = 0-4) or -O-(CH₂)_n (n = 0-4),
- V) B represents H, -SAC, -SCAC, -Ar, or -SAC-Ar,
- VI) X represents -COCH₂N₂, -COCH₂F, -COCH₂Cl, -COCH₂Br, -COCH₂I, -COCH₂OAr, -COCH₂OCOAr or -COCH₂SR¹⁷ (R¹⁷ is -SAC, -SCAC, -Ar or -SAC-Ar).
- [9] 7. The compound according to claim 1 which is selected from the following group:
 - (1) (3S)-5-[(2,6-dichlorobenzoyl)oxy] -
 - 3-({[5-methyl-3-phenyl-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-oxopentan oic acid (Iaa);
 - (2) (3S)-3-({[5-methyl-3-phenyl-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-4-oxo-5-phenoxypentanoic acid (Ibb);
 - (3) (3S)-3- $(\{[5-ethyl-3-phenyl-4,5-dihydro-5-isoxazolyl]\}$
 - carbonyl}amino)-4-oxo-5-(2,3,5,6-tetrafluorophenoxy)pentanoic acid (Icc);
 - (4) $(3S)-3-(\{[5-ethyl-3-(1-naphthyl)-4,5-dihydro-5-isoxazolyl]\}$
 - carbonyl amino)-4-oxo-5-(2,3,5,6-tetrafluorophenoxy) pentanoic acid (Idd);
 - (5) (3S)-3-({[5-ethyl-3-(2-naphthyl)-4,5-dihydro-5-isoxazolyl]
 - carbonyl amino)-4-oxo-5-(2,3,5,6-tetrafluorophenoxy) pentanoic acid (Iee);
 - (6) (3S)-3-({[5-ethyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]
 - carbonyl amino)-4-oxo-5-(2,3,5,6-tetrafhorophenoxy)pentanoic acid (Iff);
 - (7) 3-({[5-ethyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]
 - carbonyl amino)-5-fluoro-4-oxopentanoic acid (Igg);
 - (8) ethyl 3-({[5-ethyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]
 - carbonyl amino)-5-fluoro-4-oxopentanoate (Ihh);
 - (9) 5-fluoro-3-({[(5R)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-4-oxopentanoic acid (Iii);
 - (10) 3-({[5-ethyl-3-(4-quinolinyl)-4,5-dihydro-5-isoxazolyl]
 - carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Ijj);
 - (11) 3-({[3-(benzothiophen-2-yl)-5-ethyl-4,5-dihydro-5-isoxazdyl]
 - carbonyl amino)-5-fluoro-4-oxopentanoic acid (Ikk);
 - (12) (3S)-3-($\{[3-(1,3-dimethyl-1 H-indol-2-yl)-5-ethyl-4,5-dihydro-5-isoxazolyl]$

```
carbonyl}amino)-4-oxo-5-(2,3,5,6-tetrafhiorophenoxy)pentanoic acid (II);
(13) 3-({[3-(1,3-dimethyl-1 H-indol-2-yl)-5-ethyl-4,5-dihydro-5-isoxazolyl]
carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Imm);
(14) (3S)-3-([[5-ethyl-3-(1-naphthylmethyl)-4,5-dihydro-5-isoxazolyl]
carbonyl}amino)-4-oxo-5-(2,3,5,6-tetrafluorophenoxy)pentanoic acid (Inn);
(15) (3S)-5-[(2,6-dichlorobenzoyl)oxy]-3-[({5-ethyl-3-[2-(1-naphthyl)ethyl] -
4.5-dihydro-5-isoxazdyl}carbonyl)amino]-4-oxopentanoic acid (Ioo);
(16) (3S)-3-[({5-ethyl-3-[(1-naphthyloxy)methyl]} -
4,5-dihydro-5-isoxazolyl}carbonyl)amino]-4-oxo-5-(2,3,5,6-tetrafluorophenoxy)
pentanoic acid (Ipp);
(17) (3S)-3-\{[(3-\{[(4-chloro-1-naphthyl)oxy]\}
methyl}-5-ethyl-4,5-dihydro-5-isoxazolyl)carbonyl]amino}-4-oxo-5-(2,3,5,6-tetr
afluorophenoxy)pentanoic acid (Iqq);
(18) (3S,4E)-6-ethoxy-3-(\{[(5R)
)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-6-o
xo-4-hexenoic acid (Irr);
(19) (3S,4E)-3-(\{(5R)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl)
carbonyl amino)-5-(methyl sulfonyl)-4-pentenoic acid (Iss);
(20) 5-fluoro-3-(\{[(5S)-3-(1-isoquinolinyl)-5-propyl-4,5-dihydro-5-isoxazolyl\}
carbonyl}amino)-4-oxopentanoic acid (Itt);
(21) 3-([(5S)-5-ethyl-3-(1-naphthyl)-4,5-dihydro-5-isoxazolyl]
carbonyl amino)-5-fluoro-4-oxopentanoic acid (Iuu);
(22) 3-(\{[(5S)-5-\text{ethyl-}3-(2-\text{quinofinyl})-4,5-\text{dihydro-}5-\text{isoxazolyl}\})
carbonyl amino)-5-fluoro-4-oxopentanoic acid (Ivv):
(23) 3-(\{[(5R)-5-\text{ethyl-}3-(3-\text{isoquinolinyl})-4,5-\text{dihydro-}5-\text{isoxazolyl}\}
carbonyl amino)-5-fluoro-4-oxopentanoic acid (Iww);
(24) 3-({[5-ethyl-3-(8-quinolinyl)-4,5-dihydro-5-isoxazdyl]
carbonyl amino)-5-fluoro-4-oxopentanoic acid (Ixx);
(25) 3-({[5-ethyl-3-(3-quinolinyl)-4,5-dihydro-5-isoxazolyl]
carbonyl amino)-5-fluoro-4-oxopentanoic acid (Iyy);
(26) 5-fluoro-3-(\{[(5R)-5-isopropyl-3-(2-quinolinyl)-4,5-dihydro-5-isoxazolyl]
carbonyl}amino)-4-oxopentanoic acid (Izz);
(27) 3-([[5-ethyl-3-(2-isopropylphenyl)-4,5-dihydro-5-isoxazolyl]
carbonyl amino)-5-fluoro-4-oxopentanoic acid (Iaa1);
(28) 3-[({3-[3-(tert-butyl)phenyl] -
```

5-ethy I-4,5-dihydro-5-isoxazolyl carbonyl) amino]-5-fluoro-4-oxopentanoic acid (Iaa2);

 $(29) \ 3-[(\{3-[4-(tert-butyl)phenyl] -$

5-ethyl-4,5-dihydro-5-isoxazdyl}carbonyl)amino]-5-fluoro-4-oxopentanoic acid (Iaa3);

(30) 5-fluoro-3-($\{[(5R)]$

)-5-isopropyl-3-(2-isopropylphenyl)-4,5-dihydro-5-isoxazdyl]carbonyl}amino)-4-oxopentanoic acid (Iaa4);

(31) $3-[({(5R)-3-[3-(tert-butyl)phenyl]-5-isopropyl-4,5-dihydr o-$

5-isoxazolyl}carbonyl)amino]-5-fluoro-4-oxopentanoic acid (Iaa5);

(32) 3-{[(3-[1,1'-biphenyl]-3-yl-5-isopropyl-4,5-dihydro-5-isoxazolyl)carbonyl] amino}-5-fluoro-4-oxopentanoic acid (Iaa6);

(33) 3-({[5-ethyl-3-(2-pyridinyl)-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaa7);

(34) 3-[({3-[4-(tert-butyl)-2-pyridinyl] -

5-ethyl-4,5-dihydro-5-isoxazdyl}carbonyl)amino]-5-fluoro-4-oxopentanoic acid (Iaa8);

(35) $3-[({(5R)-3-[4-(tert-butyl)-2-pyridinyl]} -$

5-isopropyl-4,5-dihydro-5-isoxazdyl}carbonyl)amino]-5-fluoro-4-oxopentanoic acid (Iaa9);

(36) 3-({[5-ethyl-3-(4-isobutyl-2-pyridinyl)-4,5-dihydro -5-isoxazolyl] carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaa10);

(37) 3-({[3-(4-acetyl-2-pyridinyl)-5-ethyl-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaal 1);

(38) 3-({[3-(4-cyclopropyl-2-pyridinyl)-5-ethyl-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaa12):

(39) 3-({[3-(4-cyclopentyl-2-pyridinyl)-5-ethyl-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaal3);

(40) 3-({[(5*R*

)-3-(4-cyclopentyl-2-pyridinyl)-5-isopropyl-4,5-dihydro-5-isoxazdyl]carbonyl}a mino)-5-fluoro-4-oxopentanoic acid (Iaa14);

(41) 3-({[3-(4-cyclohexyl-2-pyridinyl)-5-ethyl-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaa15);

(42) 3-({[5-ethyl-3-(5,6,7,8-tetrahydro-1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaa16);

(43)

- 5-fluoro-3-({[5-isopropyl-3-(4-phenyl-2-pyridinyl)-4,5-dihydro-5-isoxazolyl]car bonyl}amino)-4-oxopentanoic acid (Iaa17);
- (44) (3S)-5-[(diphenylphosphoryl)oxy]-3-({[(5R)
-)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o xopentanoic acid (Iaa18);
- (45) (3*S*)-3-({[(5*R*)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-4-oxo-5-{[1-phenyl-3-(trifluoromethyl)-1 *H*-pyrazol-5-yl] oxy}pentanoic acid (Iaa19);
- (46) (3S)-5-[(4-benzyl-5-oxo-2,5-dihydro-3-furanyl)oxy-3-({[(5 R
-)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o xopentanoic acid (Iaa20);
- $(47) (3S)-5-(isobutyryloxy)-3-({[(5R)$
-)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o xopentanoic acid (Iaa21);
- (48) (3S)-3- $(\{[(5R)$ -5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl] carbonyl $\{$ amino $\}$ -4-oxo-5-hexenoic acid (Iaa22);
- (49) (3S)-3-({[(5R)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-4-oxo-5-(2-pyridinyloxy)pentanoic acid (Iaa23);
- (50) (3S)-3-({[5-ethyl-3-(2-isopropylphenyl)-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-4-oxo-5-(2-pyridinyloxy)pentanoic acid (Iaa24);
- $(51) 2-\{[(3S)-4-carboxy-3-(\{[(5R)$
-)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-2-o xobutyl]oxy}-1-methylpyridinium trifluoromethanesulfonate (Iaa25);
- $(52) 2 \{[(3S)]$
-)-4-carboxy-3-({[5-ethyl-3-(2-isopropylphenyl)-4,5-dihydro-5-isoxazolyl]carbon yl}amino)-2-oxobutyl]oxy}-1-methylpyridinium trifluoromethanesulfonate (Iaa26);
- (53) $3-(\{[3-(5-chloro-1-methyl-1 H-$
- indol-2-yl)-5-isopropyl-4,5-dihydro-5-isoxazdyl]carbonyl}amino)-5-fluoro-4-ox opentanoic acid (Iaa27);
- (54) 3-({[3-(1,5-dimethyl-1 H-indol-2-yl)-5-isopropyl-4,5-dihydro-5-isoxazolyl] carbonyl}amino)-5-fluoro-4-oxopentanoic acid (Iaa28); and
- (55) (3S)-5-fluoro-3- $(\{[(5R)$
-)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o

xopentanoic acid (Iii-1).

- [10] 8. The compound according to claim 1 which is 5-fluoro-3-({[(5 R)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o xopentanoic acid (Iii).
- [11] 9. A compound of the following formula (1a):

[13] (1a) in which

A, B, R, R¹, R², and X are defined as described in claim 1.

- [14] 10. The compound according to claim 9 which is (3.5)-5-fluoro-3-({[(5 R)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o xopentanoic acid (Iii-1).
- [15] 11. A process for preparing the compound (Iii-1) as defined in claim 10, which comprises the steps of dissolving a mixture of (3 S)-5-fluoro-3-({[(5 R) -5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o xopentanoic acid and (3R)-5-fluoro-3-({[(5 R) -5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-o xopentanoic acid in methyl t-butyl ether, adding a small amount of crystalline (3 S)-5-fluoro-3-({[(5 R)-5-isopropyl-3-(1-isoquinolinyl)-4,5-dihydro-5-isoxazolyl]carbonyl}amino)-4-oxopentanoic acid as a seed material to give a crystal, and recrystallizing this crystal from ethyl acetate/n-hexane solvent system.
- [16] 12. A therapeutic composition for preventing inflammation and apoptosis comprising the caspase inhibitor compound of formula (1), salt, or stereoisomer thereof as defined in claim 1 as an active ingredient together with the pharmaceutically acceptable carrier.
- [17] 13. The composition according to claim 12 for the treatment of dementia, cerebral stroke, brain impairment due to AIDS, diabetes, gastric ulcer, cerebral injure by hepatitis, hepatitis-induced hepatic diseases, acute hepatitis, fulminant hepatic failure, liver cirrhosis, sepsis, organ transplantation rejection, rheumatic arthritis, or cardiac cell apoptosis due to ischemic cardiac diseases.
- [18] 14. The composition according to claim 12 for the treatment of acute hepatitis or liver cirrhosis.

[25]

15. The composition according to claim 12 for the treatment of rheumatic [19] · arthritis. 16. The composition according to claim 12 which is formulated as an oral [20] preparation, an injection, or a patch. 17. The composition according to claim 12 comprising the compound (Iii) as [21] defined in claim 8 as an active ingredient. 18. The composition according to claim 12 comprising the compound (Iii-1) as [22] defined in claim 10 as an active ingredient. 19. A process for preparing the therapeutic composition for preventing in-[23] flammation and apoptosis as defined in claim 12, comprising admixing the caspase inhibitor compound of formula (1), salt, or stereoisomer thereof as defined in claim 1 with pharmaceutically acceptable carrier. 20. A method for preventing inflammation and apoptosis, comprising ad-[24] ministering an effective amount of the caspase inhibitor compound of formula (1), salt, or stereoisomer thereof as defined in claim 1 to a patient suffering from inflammation and apoptosis. 21. A use of the caspase inhibitor compound of formula (1), salt, or stereoisomer

thereof as defined in claim 1 for preventing inflammation and apoptosis.